DOCKET NO.: TIBO-0019 PATENT

Application No.: 09/530,907

Office Action Dated: June 17, 2005

This listing of claims will replace all prior versions, and listings, of claims in the application.

Listing of Claims:

1. (Currently amended) A method for screening for analytes comprising the

steps of

a) simultaneously applying a plurality of analytes to be screened onto at least one

solid support such that the analytes remain isolated from one another disposing a plurality of

analytes to be screened within individually identifiable containers such that the analytes

remain isolated from each other, wherein the individually identifiable containers are an array

of capillary tubes each of which is identifiable according to its position within the array;

b) dispensing the analytes through the open ends of the capillary tubes onto at least

one solid support to maintain the transferred contents of each container separate from those of

each other container, wherein said analytes are simultaneously applied onto the at least one

solid support;

[[b]]c) contacting said at least one analyte-carrying solid support with targets

provided in a semi-solid or liquid medium, whereby said analytes are released from the at

least one solid support to the targets, wherein each analyte when applied to the solid support

diffuses thereon so as to produce a concentration gradient; and

[[c]]d) measuring analyte-target interactions, wherein said analyte-target interactions

are measured using one or more of the following methods: microscopic, luminometric,

densitometric, isotopic, and physical measurements.

2-4. (Cancelled)

5. (Currently amended) [[A]]The method according to Claim 1 claim 1, wherein

the solid support is of a substantially flat, disc-, rectangular- or square-shape.

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6-9 (Cancelled)

10. (Currently amended) [[A]]The method according to Claim 1claim 1, wherein

the surface of the solid support onto which the analytes are applied is selected from polymers,

ceramics, metals, cellulose and glass.

11-16. (Cancelled)

17. (Currently amended) [[A]]The method according to Claim 1 wherein

the surface of the solid support is coated with a layer with molecules, [[a]] a layer with cells

or a Langmuir-Blodgett film.

18. (Currently amended) [[A]]The method according to Claim 1claim 1, wherein

the solid support is [itself] an information carrier which carries information in electronic,

magnetic or digitised digitized form.

19-23. (Cancelled)

24. (Currently amended) [[A]]The method according to Claim 1 wherein

steps a) and b) are carried out simultaneously.

25. (Cancelled)

26. (Currently amended) [[A]]The method according to Claim 1, wherein

each analyte is applied to [[a]] a rod or spherically shaped solid support.

27-28. (Cancelled)

29. (Currently amended) [[A]]The method according to Claim-Iclaim 1 wherein

the analytes are selected from chemical compounds, antigens, antibodies, DNA-probes, cells

and beads and liposomes carrying an analyte of interest.

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30. (Currently amended) [[A]]The method according to Claim 29claim 29, wherein the analytes, when applied to the solid support, are dissolved in an organic or inorganic solvent.

- 31. (Cancelled)
- 32. (Currently amended) [[A]]<u>The</u> method according to <u>Claim 1 claim 1</u> wherein the analyte is a chemical compound.
- 33. (Currently amended) [[A]]The method according to Claim 1 wherein said targets are selected from prokaryotic cells, eukaryotic cells, viruses, molecules, receptors, beads, and combinations thereof.
- 34. (Currently amended) [[A]]<u>The</u> method according to <u>Claim 33claim 33</u>, wherein the targets are cells equipped with reporter functions.
- 35. (Currently amended) [[A]]<u>The</u> method according to <u>Claim 34</u>, wherein said analyte target interactions are measurable by the effects of the analytes on the reporter functions of the cells.

36-68. (Cancelled)